DOCUMENTS/RECORDS WHICH MAY BE NEEDED FOR REVIEW DURING AN INSPECTION OF RADIATION EQUIPMENT BUREAU OF RADIATION CONTROL (BRC)

Certificate of Registration
25 TAC §289 - State Regulations - see 25 TAC §289.227(n)(3)
Operating and Safety Procedures (Individuals who are sole practitioners, sole operators, and the only occupationally exposed individual are exempt from this rule.)
Prior Notices of Violation, if applicable, documentation of corrections
Annual protective device check log (i.e, lead aprons, gloves, shields)
Equipment Performance Evaluations (this does not apply to veterinary facilities)
Verification of credentialing for each individual operating the x-ray units (this does not apply to veterinary facilities)
Personnel monitoring records (i.e., film badge reports) - (this does not apply to dental facilities)
Surveys of radiation levels in unrestricted areas - (this does not apply to dental facilities)
Records of receipt, transfer, or disposal for all x-ray units
Film processing: 25 TAC §289.227(1)
Time/temperature relationships recommended by film manufacturer
Chemical replacement - not to exceed three months
Darkroom light leak test - not to exceed six months
Safe lighting as recommended by film manufacturer
Records of any corrections or repairs (see 25 TAC §289.227 (l)(5)

ADDITIONAL RECORDS WHICH MAY BE REVIEWED FOR SPECIAL UNITS

FDA variances of machines
Current annual survey of fluoroscopic x-ray machines - (entrance exposure rate)
CT dose measurements
Radiation therapy surveys and calibrations
The most recent calibration for each linear accelerator, orthovoltage, or superficial therapy unit
Calibration of unit after change of a major component
Written spot check procedures
Initial installation survey-shielding calculations
Calibration of dosimetry system
Records, notices, and reports of therapy misadministration - see 25 TAC §289.227(bb)(2)
Licensing verification for all Physicists